# **Complete Summary**

#### **TITLE**

Diagnosis and treatment of chest pain and acute coronary syndrome (ACS): percentage of patients with acute myocardial infarction (AMI) receiving thrombolytics with a "door-to-drug time" (time from presentation to administration of drug) of less than 30 minutes.

# SOURCE(S)

Institute for Clinical Systems Improvement (ICSI). Diagnosis and treatment of chest pain and acute coronary syndrome (ACS). Bloomington (MN): Institute for Clinical Systems Improvement (ICSI); 2008 Oct. 69 p. [138 references]

#### **Measure Domain**

#### PRIMARY MEASURE DOMAIN

**Process** 

The validity of measures depends on how they are built. By examining the key building blocks of a measure, you can assess its validity for your purpose. For more information, visit the Measure Validity page.

## **SECONDARY MEASURE DOMAIN**

Does not apply to this measure

# **Brief Abstract**

# **DESCRIPTION**

This measure is used to assess the percentage of patients with acute myocardial infarction (AMI) receiving thrombolytics with a "door-to-drug time" (time from presentation to administration of drug) of less than 30 minutes.

#### **RATIONALE**

The priority aim addressed by this measure is to minimize the delay in administering thrombolytics to patients with acute myocardial infarction (AMI).

#### PRIMARY CLINICAL COMPONENT

Acute myocardial infarction (AMI); thrombolytics; "door-to-drug time" (time from presentation to administration of drug)

#### **DENOMINATOR DESCRIPTION**

Number of patients with acute myocardial infarction (AMI) receiving thrombolytics in the emergency department in the measurement period

# **NUMERATOR DESCRIPTION**

Number of patients with acute myocardial infarction (AMI) receiving thrombolytics within 30 minutes of presentation in the emergency department (All reportable "door-to-drug times"\* are rounded to the nearest minute.)

\*Formula for calculating "door-to-drug time": Time of initiation of thrombolytic therapy to patient with AMI - Time of arrival of patient with AMI in the emergency department = Door-to-drug time in minutes

# **Evidence Supporting the Measure**

# **EVIDENCE SUPPORTING THE CRITERION OF QUALITY**

 A clinical practice guideline or other peer-reviewed synthesis of the clinical evidence

#### NATIONAL GUIDELINE CLEARINGHOUSE LINK

Diagnosis and treatment of chest pain and acute coronary syndrome (ACS).

# **Evidence Supporting Need for the Measure**

#### **NEED FOR THE MEASURE**

Unspecified

## State of Use of the Measure

#### **STATE OF USE**

Current routine use

# **CURRENT USE**

Internal quality improvement

# **Application of Measure in its Current Use**

## **CARE SETTING**

Emergency Medical Services Hospitals

# PROFESSIONALS RESPONSIBLE FOR HEALTH CARE

Physicians

#### LOWEST LEVEL OF HEALTH CARE DELIVERY ADDRESSED

**Group Clinical Practices** 

# **TARGET POPULATION AGE**

Age greater than or equal to 18 years

# **TARGET POPULATION GENDER**

Either male or female

# STRATIFICATION BY VULNERABLE POPULATIONS

Unspecified

# **Characteristics of the Primary Clinical Component**

# INCIDENCE/PREVALENCE

Unspecified

# **ASSOCIATION WITH VULNERABLE POPULATIONS**

Unspecified

# **BURDEN OF ILLNESS**

Unspecified

# **UTILIZATION**

Unspecified

# **COSTS**

Unspecified

**Institute of Medicine National Healthcare Quality Report Categories** 

# **IOM CARE NEED**

Getting Better

#### **IOM DOMAIN**

Effectiveness Timeliness

#### **Data Collection for the Measure**

#### **CASE FINDING**

Users of care only

## **DESCRIPTION OF CASE FINDING**

Adults 18 and older diagnosed as having an acute myocardial infarction (AMI)

It is suggested that data collection be completed on a real-time basis. This measure references all patients to improve process sensitivity at sites where few AMI patients are routinely discharged in a given measurement period.

Should real-time data collection present insurmountable institutional obstacles, consider using the following principal diagnosis codes (International Classification of Diseases, Ninth Revision, Clinical Modification [ICD-9-CM]) for identification of patient records for abstracting:

 410 - AMI, with or without first decimal extensions in the set (0,1,2,3,4,5,6,7,8,9); as well as second decimal extensions in the set (0 or 1 only).

In addition to tracking the percentage of patients treated in less than 30 minutes, sites may choose to also track either the mean (average) or the median (middle point) of the data. Using the median is preferred. The median is the value of the middle item in the data set. The median value is preferred over the mean (average) value because it minimizes the impact of outlying data points.

For example, if one case of receiving thrombolytics took 120 minutes when the other 10 cases in the data set received them within 20 to 30 minutes, the mean would be about 34 minutes. However, the median for that same data set might be around 26 minutes, and would more accurately reflect the usual performance of the system.

Data can be collected weekly or monthly.

#### **DENOMINATOR SAMPLING FRAME**

Patients associated with provider

#### **DENOMINATOR INCLUSIONS/EXCLUSIONS**

#### **Inclusions**

Number of patients with acute myocardial infarction (AMI) receiving thrombolytics in the emergency department in the measurement period

#### **Exclusions**

Unspecified

#### RELATIONSHIP OF DENOMINATOR TO NUMERATOR

All cases in the denominator are equally eligible to appear in the numerator

# **DENOMINATOR (INDEX) EVENT**

Clinical Condition Encounter Therapeutic Intervention

#### **DENOMINATOR TIME WINDOW**

Time window is a single point in time

## **NUMERATOR INCLUSIONS/EXCLUSIONS**

#### **Inclusions**

Number of patients with acute myocardial infarction (AMI) receiving thrombolytics within 30 minutes of presentation in the emergency department (All reportable "door-to-drug times"\* are rounded to the nearest minute.)

\*Formula for calculating "door-to-drug time": Time of initiation of thrombolytic therapy to patient with AMI - Time of arrival of patient with AMI in the emergency department = Door-to-drug time in minutes

#### **Exclusions**

Unspecified

# MEASURE RESULTS UNDER CONTROL OF HEALTH CARE PROFESSIONALS, ORGANIZATIONS AND/OR POLICYMAKERS

The measure results are somewhat or substantially under the control of the health care professionals, organizations and/or policymakers to whom the measure applies.

## **NUMERATOR TIME WINDOW**

Fixed time period

#### **DATA SOURCE**

Administrative data Medical record

# **LEVEL OF DETERMINATION OF QUALITY**

Individual Case

#### PRE-EXISTING INSTRUMENT USED

Unspecified

# **Computation of the Measure**

#### **SCORING**

Rate

## **INTERPRETATION OF SCORE**

Better quality is associated with a higher score

# **ALLOWANCE FOR PATIENT FACTORS**

Unspecified

# STANDARD OF COMPARISON

Internal time comparison

# **Evaluation of Measure Properties**

# **EXTENT OF MEASURE TESTING**

Unspecified

# **Identifying Information**

# **ORIGINAL TITLE**

Percentage of patients with AMI receiving thrombolytics with a "door-to-drug time" (time from presentation to administration of drug) of less than 30 minutes.

## **MEASURE COLLECTION**

<u>Diagnosis and Treatment of Chest Pain and Acute Coronary Syndrome (ACS)</u>
Measures

# **DEVELOPER**

Institute for Clinical Systems Improvement

## **FUNDING SOURCE(S)**

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#### COMPOSITION OF THE GROUP THAT DEVELOPED THE MEASURE

Work Group Members: R. Scott Wright, MD (Work Group Leader) (Mayo Clinic) (Cardiology); Paul Spilde, PT (Park Nicollet Health Services) (Cardiac Rehabilitation); James Morrison, MD (HealthPartners Medical Group) (Cardiology); M. Danish Rizvi, MD (HealthPartners Medical Group) (Cardiology); Jackson Thatcher, MD (Park Nicollet Health Services) (Cardiology); Editha Liu, MD (Avera Health) (Hospitalist); Tonja Larson, PharmD, BCPS (Marshfield Clinic) (Pharmacy); Kathy Melsha, PharmD, BCPS (Park Nicollet Health Services) (Pharmacy); Myounghee Hanson (Institute for Clinical Systems Improvement) (Facilitator); Teresa Hunteman, MA, CPHQ (Institute for Clinical Systems Improvement) (Facilitator)

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R. Scott Wright, MD is a consultant for and receives research/grant funding from Hoffman LaRoche pertaining to clinical trial testing of Dalcetrapib.

No other work group members have potential conflicts of interest to disclose.

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#### **ADAPTATION**

Measure was not adapted from another source.

#### **RELEASE DATE**

2004 Nov

#### **REVISION DATE**

2008 Oct

#### **MEASURE STATUS**

This is the current release of the measure.

This measure updates a previous version: Institute for Clinical Systems Improvement (ICSI). Diagnosis and treatment of chest pain and acute coronary syndrome (ACS). Bloomington (MN): Institute for Clinical Systems Improvement (ICSI); 2006 Oct. 76 p.

## SOURCE(S)

Institute for Clinical Systems Improvement (ICSI). Diagnosis and treatment of chest pain and acute coronary syndrome (ACS). Bloomington (MN): Institute for Clinical Systems Improvement (ICSI); 2008 Oct. 69 p. [138 references]

#### **MEASURE AVAILABILITY**

The individual measure,"Percentage of Patients with AMI Receiving Thrombolytics with a "Door-to-Drug Time" (Time from Presentation to Administration of Drug) of Less Than 30 Minutes," is published in "Health Care Guideline: Diagnosis and Treatment of Chest Pain and Acute Coronary Syndrome (ACS)." This document is available from the Institute for Clinical Systems Improvement (ICSI) Web site.

For more information, contact ICSI at, 8009 34th Avenue South, Suite 1200, Bloomington, MN 55425; phone: 952-814-7060; fax: 952-858-9675; Web site: <a href="https://www.icsi.org">www.icsi.org</a>; e-mail: <a href="https://icsi.info@icsi.org">icsi.info@icsi.org</a>.

# **NQMC STATUS**

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